Application No.: 10/084,621

Response to Final Office Action dated August 13, 2003

Response filed November 13, 2003

#### Amendments to the Claims:

This listing of claims will replace all prior versions, and listings, of claims in the application:

# **Listing of Claims**:

## Claim 1. (Currently amended):

A transgenic mouse whose genome contains a homozygous disruption of both the endogenous Gpx1 gene and Gpx2 gene wherein said mouse develops cancer.

## Claim 2. (Previously presented):

A cell from the transgenic mouse of claim 1.

## Claims 3-4. (Canceled).

## Claim5. (Previously presented):

A cell of claim 2 which is selected from the group consisting of stem cells, epithelial cells and myelofibroblasts.

## Claim 6. (Previously presented):

The transgenic mouse of claim 1 wherein the genetic background of the mouse is selected from the group consisting of a B6 mouse, a 129Sv/J hybrid mouse, a 129S3 hybrid mouse and a ½ B6, 1/4 129Sv/J and 1/4 129S3 hybrid mouse.

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Claim 7. (Previously presented):

A transgenic mouse as in claim 1 which further comprises a mouse which is a germ free mouse.

Claims 8-39 canceled.

Claim 40. (Currently amended):

A transgenic double knockout mouse <u>as in claim 1</u> whose genome comprises a homozygous disruption of the endogenous *Gpx1* gene and a homozygous disruption of the endogenous *Gpx2* gene, wherein each disruption comprises the insertion of a transgene, and wherein the combined disruptions result in a decreased level of GPX-1 and GPX-GI production and decreased number of cells producing GPX-I and GPX-GI in the transgenic mouse as compared to a nontransgenic mouse.

Claim 41. (Currently amended):

A transgenic double knockout mouse as in claim 40 1 which exhibits one or more physiological symptoms selected from the group consisting of ileitis, colitis, hypothermia, decreased rate of weight gain, perianal ulceration, diarrhea, wasting syndrome, inflammatory bowel disease and cancer of the lower gastro-intestinal tract, one or more tumors in the small bowel and cancer of the small intestine.

Claim 42. (Previously presented):

A cell isolated from a double knockout mouse as in claim 40.

Application No.: 10/084,621

Response to Final Office Action dated August 13, 2003

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#### Claim 43. (Previously presented):

A cell as in claim 42, selected from the group consisting of a stem cell, an epithelial cell and a myofibroblast.

## Claim 44. (Previously presented):

A cell as in claim 43 which is a stem cell.

## Claim 45. (Previously presented):

A cell as in claim 43 which is an epithelial cell.

### Claim 46. (Previously presented):

A cell as in claim 43 which is a myofibroblast.

## Claim 47. (Previously presented):

A transgenic double knockout mouse as in claim 40 which further comprises a mouse which is a germ free mouse.

#### Claim 48. (Previously presented):

A transgenic double knockout mouse as in claim 1 wherein said knockout mouse is a mouse with a B6 genetic background.

## Claim 49. (Currently amended):

A transgenic double knockout mouse as in claim 1 wherein said knockout mouse is a mouse with a hybrid mouse having a ½ B6, ¼ 129 SuJ and ¼ 129 S3 genetic background.

Claims 50-66 canceled.